

(19)



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(11)

**EP 0 874 984 B1**

(12)

**EUROPEAN PATENT SPECIFICATION**

(45) Date of publication and mention  
of the grant of the patent:  
28.11.2001 Bulletin 2001/48

(51) Int Cl.7: G01N 27/327, C12Q 1/00

(21) Application number: 96922651.3

(86) International application number:  
PCT/US96/11240

(22) Date of filing: 28.06.1996

(87) International publication number:  
WO 97/02487 (23.01.1997 Gazette 1997/05)

(54) **ELECTROCHEMICAL BIOSENSOR TEST STRIP**

**TESTSTREIFEN FÜR EINEN ELEKTROCHEMISCHEN BIOSENSOR**

**BANDE D'ESSAI DE BIOCAPTEUR ELECTROCHIMIQUE**

(84) Designated Contracting States:  
DE ES FR GB IT

(30) Priority: 30.06.1995 US 496939

(43) Date of publication of application:  
04.11.1998 Bulletin 1998/45

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**EP 0 874 984 B1**

Description

Field of The Invention

5 [0001] This invention relates generally to the determination of the concentration of analytes in fluids and more specifically to an amperometric biosensor for use in such determination.

Background of The Invention

10 [0002] Biosensors are not new. Their use in the determination of concentrations of various analytes in fluids is also known.

[0003] Nankai et al., WO 86/07632, published December 31, 1986, discloses an amperometric biosensor system in which a fluid containing glucose is contacted with glucose oxidase and potassium ferricyanide. The glucose is oxidized and the ferricyanide is reduced to ferrocyanide. (This reaction is catalyzed by glucose oxidase.) After two minutes, an electrical potential is applied and a current caused by the re-oxidation of the ferrocyanide to ferricyanide is obtained. The current value, obtained a few seconds after the potential is applied, correlates to the concentration of glucose in the fluid.

[0004] Because Nankai et al. discloses a method in which the reaction of glucose and ferricyanide may run to completion prior to the application of an electrical potential, this method is referred to as the "end-point" method of amperometric determination.

[0005] Nankai et al. discloses a system, wherein the glucose oxidase and potassium ferricyanide are held on a nonwoven nylon mesh. The mesh is positioned so that it is in contact with a working electrode, a counter electrode and a reference electrode. The total surface area of the counter and reference electrodes is twice that of the working electrode.

25 [0006] Wogoman, EP 0 206 218, published Dec. 30, 1986 discloses a biosensor having two electrodes, the electrodes being made of different electrically conducting materials. For example, the anode is formed from an anode material, such as platinum, and the cathode is formed from a cathode material, such as silver. The anode is coated with an enzyme. In a preferred embodiment, the coated electrode is covered with an elastomer that is permeable to glucose.

[0007] Pottgen et al., WO 89/08713, published Sept. 21, 1989, discloses the use of a two electrode biosensor, wherein the electrodes are made of the same noble metal, but one of the electrodes (referred to as a pseudoreference electrode) is larger than the other (working) electrode.

[0008] Recently, Pollmann et al., U.S. Patent No. 5,288,636, issued Feb. 22, 1994, disclosed an electrochemical biosensor test strip that includes working and counter electrodes of substantially the same size and made of the same electrically conducting materials. The Pollmann et al. test strip includes a reagent well that will accommodate a testing sample of human whole blood from about 10 to about 70 microliters. However, below about 13 microliters, errors in the measurement of an analyte, such as glucose, from a whole blood sample may result (low dosing errors). Generally, the low dosing error is manifested as an understated measurement of the analyte, or no measurement of the analyte by the meter used in conjunction with the test strip. Low dosing errors are a particular concern for infants and elderly persons who often have difficulty in expressing a reasonably sized blood drop for testing upon pricking their finger with a lancet.

40 [0009] Accordingly, it is highly desirable to design a test strip that requires a minimum volume of blood for the testing of an analyte, such as blood glucose.

Summary of the Invention

45 [0010] The invention is an electrochemical biosensor test strip that has a lower minimum volume blood sample requirement than prior art strips of similar construction. The present inventive test strip has a smaller reagent well and smaller spreading mesh than similar prior art strips. Further, the reagent well is positioned differently than in similar prior art test strips. The minimum blood volume sample requirement for the new strip is about 9 microliters.

50 [0011] The smaller sample volume requirement means fewer low sample volume dosing errors result when measuring an analyte, such as glucose, from a whole blood sample. This result is especially important for those persons, such as infants and the elderly, who have difficulty expressing a reasonably sized drop of blood by pricking their finger with a lancet. Also, with the present inventive strip it is easier for the meter, which collects current measurements and correlates those measurements to a concentration of analyte from a sample, to discriminate low sample volume dosing errors. Further, the smaller reagent well requires less reagent per biosensor strip, thereby increasing the production volume for mass production of biosensor test strips.

55 [0012] Additionally, when the spreading mesh is affixed to the test strip by an adhesive tape, the tape includes a hole that exposes the reagent well and spreading mesh, and further includes air vents on opposing sides of the hole. These

air vents reduce the occurrence of air bubbles trapped in the reagent well when a sample is being tested. Air bubbles can produce testing errors.

# Brief Description of the Drawings

[0013]

FIG. 1 is an exploded view of the present inventive biosensor test strip.

FIG. 2 is a top view of the biosensor test strip without the reagent, spreading mesh, and adhesive tape with air vents.

FIG. 3 is a top view of the fully constructed, preferred biosensor test strip.

FIG. 4 is a cross-sectional view of the biosensor of FIG. 3 along lines 21-21.

FIG. 5 illustrates hypothetical calibration curves for different lots of biosensor test strips.

## Description of the Preferred Embodiment

[0014] The present inventive biosensor test strip is similar to the preferred embodiment of the test strip described in Pollmann et al., U.S. Patent No. 5,288,636, issued Feb. 22, 1994. However, the Pollmann et al. strip has a construction such that too many low dosing errors result when whole blood samples below about 13 microliters are tested for blood glucose.

[0015] In the present inventive test strip, reagent well 9 (Fig. 4) has been reduced in size over the Pollmann et al. reagent well and repositioned so that a smaller surface area of the counter electrode 5 than the working electrode 4 is exposed by cutout portion 8, which forms reagent well 9. (Figs. 1-4) Mesh 13, which is a spreading mesh, is also reduced in size over the Pollmann et al. mesh. (Figs. 1, 3, 4) These changes in strip architecture result in a test strip that can accurately measure an analyte, such as glucose, from a minimum whole blood sample of about 9 microliters.

[0016] Referring specifically to Figs. 1 through 4, there is shown the presently preferred embodiment of the inventive biosensor test strip.

[0017] Test strip 1 comprises first and second electrically insulating layers 2 and 3, respectively. Any useful insulating material will be suitable. Typically, plastics, such as vinyl polymers and polyimides provide the electrical and structural properties which are desired. Preferably, these layers are Melinex® 329, 7 mil.

[0018] The biosensor test strip shown in Figs. 1 through 4 is intended to be mass produced from rolls of material, necessitating the selection of a material which is sufficiently flexible for roll processing and at the same time sufficiently stiff to give a useful stiffness to the finished biosensor test strip.

[0019] Layers 2 and 3 may be of any useful thickness. In a preferred embodiment, layers 2 and 3 are about 180 µm (7 mil) thick.

[0020] Working electrode 4 and counter electrode 5 are preferably deposited on a backing of insulator material 7, such as polyimide, to reduce the possibility of tearing the electrode before it is affixed to layer 2. Working electrode 4 and counter electrode 5 are substantially the same size and are made of the same electrically conducting material. Examples of electrically conducting materials that may be used are palladium, platinum, gold, silver, carbon, titanium, and copper. Noble metals are preferred because they provide a more constant, reproducible electrode surface area. Palladium is particularly preferred because it is one of the more difficult noble metals to oxidize and because it is a relatively inexpensive noble metal. Silver is not preferred because it is more readily oxidized by air than the other noble metals listed above. Preferably, electrodes 4 and 5 are about 0.1 µm (micron) thick and backing 7 is about 25 µm (microns) thick (commercially available from Courtaulds Performance Films in California and Southwall Technologies, Inc.).

[0021] Electrodes 4 and 5 must be sufficiently separated so that the electrochemical events at one electrode do not interfere with the electrochemical events at the other electrode. The preferred distance between electrodes 4 and 5 is about 1.2 millimeters.

[0022] In the preferred embodiment, electrodes 4 and 5, affixed to backing 7, are unspooled from reels and attached to layer 2 by the use of hot melt adhesive (not shown). Electrodes 4 and 5 also preferably extend from one end of layer 2 to the other end in parallel configuration.

[0023] Insulating layer 3 is fixed on top of layer 2 and electrodes 4 and 5 by the use of hot melt adhesive (not shown). Layer 3 includes cutout portion 8, which defines reagent well 9. Both the size and the position of cutout portion 8 are critical to the invention. Cutout portion 8 must be sufficiently small and must be sufficiently positioned such that in combination with the spreading mesh, described below, a minimum whole blood sample volume of about 9 microliters

may be accurately analyzed by the test strip. The preferred size of cutout portion 8 is 4 millimeters by 4.2 millimeters.

[0024] In the preferred embodiment, the 4 mm side of cutout portion 8 runs parallel to the long side of the test strip shown in Figs. 1-4. Importantly, cutout portion 8 is positioned over electrodes 4 and 5 such that a smaller surface area of counter electrode 5 than working electrode 4 is exposed. Preferably, the exposed surface area of working electrode 4 is twice as large as the exposed surface area of counter electrode 5. Surprisingly, offsetting cutout portion 8 to expose a smaller surface area for the counter electrode than the working electrode does not adversely affect measurement of an analyte from a sample being measured. In this preferred embodiment, electrodes 4 and 5 are 1.5 mm in width.

[0025] Biosensor test strip 1 may be accompanied by a power source (not shown) in a electrical connection with the working and counter electrodes and a current measuring meter (not shown) which is also in a electrical connection with the working and counter electrodes.

[0026] Biosensor reagent 11 (Fig. 4) is placed in well 9 so that it covers substantially all of exposed surfaces 10 and 20 of working electrode 4 and counter 5, respectively. (Figs. 2-4) An example of a reagent that may be used in the biosensor test strip of the present invention is a reagent for measuring glucose from a whole blood sample.

[0027] A protocol for making a glucose reagent utilizing the enzyme glucose oxidase and ferricyanide as the oxidized form of the redox mediator is as follows:

Step 1- Prepare 1 liter (in a volumetric flask) of a buffer/NATROSOL® mixture by adding 1.2000 grams (g) NATROSOL® -250 M to 0.740 M aqueous potassium phosphate buffer (including 80.062 g monobasic potassium phosphate and 26.423 g dibasic potassium phosphate) at pH 6.25. Allow the NATROSOL® to stir and swell for 3 hours.

Step 2- Prepare an AVICEL® mixture by stirring 14.0000 g AVICEL® RC-591 F and 504.7750 g water for 20 minutes.

Step 3- Prepare a TRITON® mixture by adding 0.5000 g TRITON® X-100 to 514.6000 g of the buffer/ NATROSOL® mixture and stir for 15 minutes.

Step 4- While stirring, add the total TRITON® mixture dropwise with an addition funnel or buret to the total AVICEL® mixture. Once addition is complete, continue stirring overnight.

Step 5- To the mixture resulting from Step 4, add, while stirring, 98.7750 g potassium ferricyanide. (Add a little potassium ferricyanide at a time to allow the potassium ferricyanide to dissolve as added.)

Step 6- Stir the resulting mixture of Step 5 for 20 minutes.

Step 7- Adjust the pH of the mixture resulting from Step 6 to 6.25 by adding potassium hydroxide.

Step 8- To the resulting mixture of Step 7, add 9.1533 g glucose oxidase (218.50 tetramethyl benzidine units per milligram (mg) from Biozyme) and stir at least 20 minutes.

Step 9- To the resulting mixture of Step 8, add 20 g potassium glutamate and stir at least 20 minutes.

Step 10- Filter the resulting mixture of Step 9 through a 100 µm (micron) sieve bag to remove any AVICEL® clumping. The filtrate is the resulting reagent composition (reagent 11), which is added to reagent well 9 and is then dried at about 50 °C for about 3 minutes.

[0028] In the preferred embodiment for glucose determination, 4 microliters of reagent made by the above-stated protocol is added to well 9 formed by cutout 8. This amount of reagent 11 will substantially cover surface areas 10 and 20 of the electrodes 4 and 5 (Fig. 2) and will also contain a sufficient amount of ferricyanide, and a sufficient amount of enzyme (glucose oxidase) to catalyze the oxidation of glucose (from a sample of human whole blood) and the reduction of ferricyanide to completion, as defined herein, within about 20 seconds. (Prior to adding the reagent to well 9, it is preferable to treat well 9 with a 600 Watt corona arc, gapped at 1/40,000 inch on a processing line travelling at 4 meters per minute, to make well 9 more hydrophilic, thereby allowing the reagent to spread more evenly in the well.)

[0029] Another glucose reagent that may be formulated includes 300 millimolar potassium ferricyanide, 250 millimolar potassium phosphate buffer, 14 grams microcrystalline cellulose (AVICEL® RC-591 F) per liter of reagent, 0.6 grams hydroxyethylcellulose (NATROSOL® -250 M) per liter of reagent, 0.5 grams Triton® X-100 surfactant per liter of reagent, 37 millimolar sodium succinate, and 1.57 million tetramethyl benzidine units of glucose oxidase per liter of reagent. Sodium hydroxide (6 Normal solution) is used to titrate this reagent to a pH of 6.6. This reagent may be formulated by



the same protocol described above, but amounts of components should be adjusted and components substituted (sodium succinate for potassium glutamate and sodium hydroxide for potassium hydroxide) to achieve the component concentrations stated above. Drying of this reagent in reagent well 9 typically results in a loss of enzyme activity of about 30-35%.

[0030] After drying reagent 11, a spreading mesh 13, which has been impregnated with a surfactant, is placed over cutout portion 8 and is affixed to second electrical insulator 3. Spreading mesh 13 is preferably a polyester monofilament mesh from ZBF (Zurich Bolting Cloth Mfg. Co. Ltd., Rüschlikon, Switzerland). The spreading mesh is preferably dipped in a solution of 0.8% (wt.:vol.) dioctylsodium sulfosuccinate (DONS) in a solution of 50:50 (vol.:vol.) methanol:water, and then dried. Spreading mesh 13 must be small enough such that in combination with the size of cutout portion 8 and placement of cutout portion 8 the biosensor strip will accurately measure analyte from a minimum whole blood sample of about 9 microliters. The preferable dimensions of spreading mesh 13 are 6 mm x 5.8 mm. In the most preferred biosensor strip, the 6 mm side of the mesh is parallel to the long side of the strip shown in Figs. 1-4.

[0031] Preferably, spreading mesh 13 is affixed to adhesive tape 14, which includes hole 15. (Figs. 1, 3, 4) Adhesive tape 14 is preferably made of polyester with an adhesive backing. (Available from Tapemark, Medical Products Division, 223 E. Marie Ave., St. Paul, Minnesota 55118) Adhesive tape 14 is preferably dyed maroon and hole 15 provides a target area for application of a sample to be analyzed by the biosensor. Hole 15 exposes at least a portion of spreading mesh 13 and cutout portion 8, and preferably exposes substantially all of cutout portion 8. Tape 14 preferably includes slits 16, as shown in Figs. 1 and 3, located on opposing sides of hole 15. (Two slits 16 are shown in Figs. 1 and 3, but one slit may be sufficient.) Slits 16 constitute air vents, which reduce the occurrence of air bubbles trapped in the reagent well upon the addition of a sample such whole blood to the reagent well. Reducing the occurrence of air bubbles trapped in reagent well 9 results in fewer testing errors.

[0032] After drying the reagent and affixing the spreading mesh, the roll-formed biosensors are separated by die punching to form discrete biosensors, which are used in conjunction with 1) a power source in electrical connection with the working and counter electrodes and capable of supplying an electrical potential difference between the working and counter electrodes sufficient to cause diffusion limited electrooxidation of the reduced form of the redox mediator at the surface of the working electrode, and 2) a meter in electrical connection with the working and counter electrodes and capable of measuring the diffusion limited current produced by oxidation of the reduced form of the redox mediator when the above-stated electrical potential difference is applied.

[0033] The meter described above will normally be adapted to apply an algorithm (discussed below) to the current measurement, whereby an analyte concentration is provided and visually displayed. Improvements in such power source, meter, and biosensor system are the subject of commonly assigned U.S. Patent Number 4,963,814, issued October 16, 1990; U.S. Patent No. 4,999,632, issued March 12, 1991; U.S. Patent No. 4,999,582, issued March 12, 1991; U.S. Patent No. 5,243,516, issued September 7, 1993; U.S. Patent No. 5,352,351, issued Oct. 4, 1994; U.S. Patent No. 5,366,609, issued Nov. 22, 1994; White et al., U.S. Patent Application Serial No. 08/073,179, filed 6/8/93 (Issue Fee mailed 12/27/94), now U.S. Patent No. 5,405,511, issued April 11, 1995; and White et al., U.S. Patent Application Serial No. 08/343,363, filed 11/22/94 (Issue Fee mailed 5/5/95), now U.S. Patent No. 5,438,271, issued August 1, 1995.

[0034] For easy electrical connection of the power source and meter, additional cutout portion 12 (Figs. 1 through 4), exposing portions of the working and counter electrodes, are preferably provided in the biosensor device.

[0035] The biosensor device described above may be used to determine the concentration of an analyte in a fluid sample by performing the following steps:

a) contacting a fluid sample, such as whole blood, with a reagent (described above) that substantially covers surface areas 10 and 20 of working and counter electrodes 4 and 5, respectively;

b) allowing the reaction between the analyte and the oxidized form of the redox mediator to go to completion, as defined herein;

c) subsequently applying a potential difference between the electrodes sufficient to cause diffusion limited electrooxidation of the reduced form of the redox mediator at the surface of the working electrode;

d) thereafter measuring the resulting diffusion limited current; and

e) correlating the current measurement to the concentration of analyte in the fluid. (Reaction completion is defined as sufficient reaction between the analyte and the oxidized form of the redox mediator to correlate analyte concentration to diffusion limited current generated by oxidation of the reduced form of the redox mediator at the surface of the working electrode.)

[0036] Many analyte-containing fluids may be analyzed. For example, analytes in human body fluids such as whole blood, blood serum, urine and cerebrospinal fluid may be measured. Also, analytes found in fermentation products and in environmental substances, which potentially contain environmental contaminants, may be measured.

[0037] When measuring analytes found in human body fluids, especially whole blood, the potential difference applied between the electrodes is preferably no more than about 500 millivolts. When a potential difference above about 500 millivolts is applied between the electrodes, oxidation of the working electrode surface (for palladium) and of some blood components may become intolerable, thereby preventing an accurate and precise correlation of current to analyte concentration. For an assay of glucose in a whole blood sample, wherein the oxidized form of the redox mediator is ferricyanide, a potential difference from about 150 millivolts to about 500 millivolts may be applied between the electrodes to achieve diffusion limited electrooxidation of the reduced form of the redox mediator at the surface of the working electrode. Preferably, about 300 millivolts potential difference is applied between the electrodes.

[0038] Current generated from the oxidation of the reduced form of the redox mediator may be measured at any time from about 0.5 seconds to about 30 seconds after the potential difference is applied between the electrodes. At less than about 0.5 seconds, diffusion limited current is difficult to measure due to the charging current. After about 30 seconds, convection becomes significant, thereby interfering with the measurement of a diffusion limited current.

[0039] The current measured during the assay of an analyte from a fluid sample may be correlated to concentration of the analyte in the sample by application of an algorithm by the current measuring meter. The algorithm may be a simple one, as illustrated by the following example:

$$[\text{Analyte}] = C i_{7.5} + d$$

wherein [Analyte] represents the concentration of the analyte in the sample (see Fig. 5),  $i_{7.5}$  is the current (in microamps) measured at 7.5 seconds after application of the potential difference applied between the electrodes, C is the slope of line 22 (Fig. 5), and d is the axis intercept (Fig. 5).

[0040] By making measurements with known concentrations of analyte, calibration curve 22 (Fig. 5) may be constructed. This calibration will be stored in the Read Only Memory (ROM) key of the meter and will be applicable to a particular lot of biosensor test strips. Lines 24 and 26 in Fig. 5 represent other hypothetical calibration curves for two other different lots of biosensor test strips. Calibration for these biosensor lots would generate slightly different values for C and d in the above algorithm.

[0041] In analysis of glucose from a sample of human whole blood, 20  $\mu$ l of whole blood is preferably added to the above-stated glucose reagent. The reaction of glucose and ferricyanide is allowed to go to completion, thereby forming gluconic acid and ferrocyanide. This reaction normally requires a short time, preferably less than about 20 seconds, to go to completion. About twenty seconds after addition of the whole blood sample, a potential difference of about 300 millivolts is applied between the electrodes, thereby oxidizing ferrocyanide to ferricyanide at the surface of the working electrode. Current measurements are made at 0.5 second intervals from 1 second to 7.5 seconds after the potential difference is applied between the electrodes. These current measurements are correlated to the concentration of glucose in the blood sample.

[0042] In this example of measuring glucose from a blood sample, current measurements are made at different times (from 1 second to 7.5 seconds after application of the potential difference), rather than at a single fixed time (as described above), and the resulting algorithm is more complex and may be represented by the following equation:

$$[\text{Glucose}] = C_1 i_1 + C_2 i_2 + C_3 i_3 + \dots C_n i_n + d,$$

wherein  $i_1$  is the current measured at the first measurement time (1 second after application of the 300 millivolt potential difference),  $i_2$  is the current measured at the second measurement time (1.5 seconds after application of the 300 millivolt potential difference),  $i_3$  is the current measured at the third measurement time (2 seconds after application of the 300 millivolt potential difference),  $i_n$  is the current measured at the  $n^{\text{th}}$  measurement time (in this example, at the 14<sup>th</sup> measurement time or 7.5 seconds after application of the 300 millivolt potential difference),  $C_1$ ,  $C_2$ ,  $C_3$ , and  $C_n$  are coefficients derived from a multivariate regression analysis technique, such as Principle Components Analysis or Partial Least Squares, and d is the regression intercept (in glucose concentration units). (A modification of this procedure may be used in the event that calibration curves illustrated by Fig. 5 have considerable curvature.)

[0043] Alternatively, the concentration of glucose in the sample being measured may be determined by integrating the curve generated by plotting current, i, versus measurement time over some time interval (for example, from 1 second to 7.5 seconds after application of the 300 millivolt potential difference), thereby obtaining the total charge transferred during the measurement period. The total charge transferred is directly proportional to the concentration of glucose in the sample being measured.

[0044] Further, the glucose concentration measurement may be corrected for differences between environmental temperature at the time of actual measurement and the environmental temperature at the time calibration was performed. For example, if the calibration curve for glucose measurement was constructed at an environmental temperature of 23°C, the glucose measurement is corrected by using the following equation:

$$[\text{Glucose}]_{\text{corrected}} = [\text{Glucose}]_{\text{measured}} \times (1 - K(T - 23^\circ\text{C})),$$

wherein T is the environmental temperature (in °C) at the time of the sample measurement and K is a constant derived from the following regression equation:

$$Y = K(T - 23),$$

wherein

$$Y = \frac{[\text{Glucose}]_{\text{measured at } 23^\circ\text{C}} - [\text{Glucose}]_{\text{measured at } T^\circ\text{C}}}{[\text{Glucose}]_{\text{measured at } T^\circ\text{C}}}$$

In order to calculate the value of K, each of a multiplicity of glucose concentrations is measured by the meter at various temperatures, T, and at 23°C (the base case). Next, a linear regression of Y on T-23 is performed. The value of K is the slope of this regression.

[0045] The glucose concentration of a sample may be accurately and precisely measured by the present inventive method utilizing the present inventive biosensor. Further, when a sample of human whole blood is measured, error due to hematocrit effect is insignificant in the range of 30-55 % hematocrit. Other examples of enzymes and redox mediators (oxidized form) that may be used in measuring particular analytes by the present invention are listed below in Table 1.

TABLE 1

ANALYTE	ENZYMES	REDOX MEDIATOR (OXIDIZED FORM)	ADDITIONAL MEDIATOR
GLUCOSE	GLUCOSE DEHYDROGENASE AND DIAPHORASE	FERRICYANIDE	
GLUCOSE	GLUCOSE-DEHYDROGENASE (QUINOPROTEIN)	FERRICYANIDE	
CHOLESTEROL	CHOLESTEROL ESTERASE AND CHOLESTEROL OXIDASE	FERRICYANIDE	2,6-DIMETHYL-1,4-BENZOQUINONE 2,5-DICHLORO-1,4-BENZOQUINONE OR PHENAZINE ETHOSULFATE
HDL CHOLESTEROL	CHOLESTEROL ESTERASE AND CHOLESTEROL OXIDASE	FERRICYANIDE	2,6-DIMETHYL-1,4-BENZOQUINONE 2,5-DICHLORO-1,4-BENZOQUINONE OR PHENAZINE ETHOSULFATE
TRIGLYCERIDES	LIPOPROTEIN LIPASE, GLYCEROL KINASE, AND GLYCEROL-3-PHOSPHATE OXIDASE	FERRICYANIDE OR PHENAZINE ETHOSULFATE	PHENAZINE METHOSULFATE

TABLE 1 (continued)

ANALYTE	ENZYMES	REDOX MEDIATOR (OXIDIZED FORM)	ADDITIONAL MEDIATOR
LACTATE	LACTATE OXIDASE	FERRICYANIDE	2,6-DICHLORO- 1,4-BENZOQUINONE
LACTATE	LACTATE DEHYDROGENASE AND DIAPHORASE	FERRICYANIDE, PHENAZINE ETHOSULFATE, OR PHENAZINE METHOSULFATE	
LACTATE DEHYDROGENASE	DIAPHORASE	FERRICYANIDE, PHENAZINE ETHOSULFATE, OR PHENAZINE METHOSULFATE	
PYRUVATE	PYRUVATE OXIDASE	FERRICYANIDE	
ALCOHOL	ALCOHOL OXIDASE	PHENYLENEDIAMINE	
BILIRUBIN	BILIRUBIN OXIDASE	1-METHOXY- PHENAZINE METHOSULFATE	
URIC ACID	URICASE	FERRICYANIDE	

[0046] In some of the examples shown in Table 1, at least one additional enzyme is used as a reaction catalyst. Also, some of the examples shown in Table 1 may utilize an additional mediator, which facilitates electron transfer to the oxidized form of the redox mediator. The additional mediator may be provided to the reagent in lesser amount than the oxidized form of the redox mediator.

[0047] When compared to the preferred embodiment of the closest prior art biosensor test strip, disclosed in Pollmann et al., the present inventive biosensor has the following distinguishing features:

1. reagent well 9 is 30% smaller;
2. when the working and counter electrodes are substantially the same size, the exposed surface area of the counter electrode in the reagent well is less than the exposed surface area of the working electrode in the reagent well;
3. a smaller amount of reagent is needed in the reagent well (4 microliters of reagent vs. 6 microliters of reagent in the preferred embodiment of Pollmann et al.);
4. a smaller spreading mesh is needed; and
5. air vents are included on opposing sides of the reagent well.

[0048] A smaller sample volume requirement to properly dose the test strip means fewer underdosing errors will result. This result is especially important for those persons, such as infants and the elderly who have difficulty in obtaining a reasonably sized blood drop after pricking their finger with a lancet. The present inventive strip makes it easier for a current measuring meter to discriminate low sample volume dosing errors. Also, using less reagent per sensor increases production volume for mass producing sensors. Further, providing side air vents near the reagent well reduces the occurrence of air bubbles trapped in the reagent well, which results in fewer testing errors.

[0049] The present invention has been disclosed in the above teachings and drawings with sufficient clarity and conciseness to enable one skilled in the art to make and use the invention, to know the best mode for carrying out the invention, and to distinguish it from other inventions and what is old. Many inventions and obvious adaptations of the invention will readily come to mind, and these are intended to be contained within the scope of the invention as claimed herein.



Claims

1. A device (1) for detecting or measuring the concentration of an analyte, comprising:

5 a first electrical insulator (2);  
a pair of electrodes (4, 5) consisting of working and counter electrodes of substantially the same size, the electrodes (4, 5) being made of the same electrically conducting materials and being supported on the first electrical insulator (2);  
10 a second electrical insulator (3), overlaying the first electrical insulator (2) and the electrodes (4, 5) and including a cutout portion (8);  
a reagent (11) for detecting or measuring the concentration of the analyte, the reagent (11) substantially covering the exposed electrode surfaces in the cutout portion (8) and being in sufficient amount for analyzing the analyte in a whole blood sample of at least about 9 microliters and  
15 a spreading mesh (13), overlaying the cutout portion (8) and affixed to the second electrical insulator (3),

characterized in that

20 the cutout portion (8) exposes a smaller surface area of the counter electrode than the working electrode;  
the spreading mesh (13) is impregnated with a surfactant;  
the cutout portion (8) and spreading mesh (13) are of sufficient size to receive a whole blood sample of at least about 9 microliters for analyzing the analyte, and  
the spreading mesh (13) is affixed to the second electrical insulator (3) by tape (14) having an adhesive on one side and a hole (15) that exposes at least a portion of the spreading mesh (13) and the cutout portion (8),  
25 and wherein the tape (14) also includes at least one slit (16) near the hole (15), thereby providing at least one air vent.

2. The device of claim 1, wherein the tape (14) includes slits (16) on opposing sides of the hole (15), thereby providing two air vents.
- 30 3. The device of claim 1, wherein the cutout portion (8) is 4 millimeters by 4.2 millimeters.
4. The device of claim 3, wherein the spreading mesh (13) is 6 millimeters by 5.8 millimeters.
5. The device of claim 4, wherein the spreading mesh (13) is impregnated with dioctylsodium sulfosuccinate.
- 35 6. The device of claim 5, wherein the hole (15) in the tape (14) exposes substantially all of the cutout portion (8).
7. The device of claim 1 or 6, further comprising a current measuring meter in electrical connection with the working and counter electrodes.
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Patentansprüche

1. Vorrichtung (1) zum Erfassen oder Messen der Konzentration eines Analyten, umfassend:

45 einen ersten elektrischen Isolator (2);  
ein Elektrodenpaar (4, 5), bestehend aus Arbeits- und Gegenelektrode von im wesentlichen gleicher Größe, wobei die Elektroden (4, 5) aus den gleichen elektrisch leitenden Materialien gefertigt sind und auf dem ersten elektrischen Isolator (2) gehalten werden;  
50 einen zweiten elektrischen Isolator (3), der über dem ersten elektrischen Isolator (2) und den Elektroden (4, 5) liegt und einen Ausschnitt (8) umfaßt;  
ein Reagens (11) zum Erfassen oder Messen der Konzentration des Analyten, wobei das Reagens (11) im wesentlichen die freiliegenden Elektrodenoberflächen im Ausschnitt (8) bedeckt und in ausreichender Menge vorliegt, um in einer Vollblutprobe von wenigstens etwa 9 µl den Analyten zu analysieren; und  
55 ein Verteilungsnetz (13), das über dem Ausschnitt (8) liegt und am zweiten elektrischen Isolator (3) befestigt ist.

dadurch gekennzeichnet, daß

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der Ausschnittteil (8) eine kleinere Oberfläche an der Gegenelektrode als an der Arbeitselektrode freiläßt, das Verteilungsnetz (13) mit einem Tensid imprägniert ist, der Ausschnittteil (8) und das Verteilungsnetz (13) von hinreichender Größe sind, um eine Vollblutprobe von wenigstens etwa 9 µl zur Analyse des Analyten aufzunehmen, und das Verteilungsnetz (13) am zweiten elektrischen Isolator (3) mit einem Band (14) befestigt ist, das ein Haftmittel auf der einen Seite sowie ein Loch (15) aufweist, das wenigstens einen Teil des Verteilungsnetzes (13) und des Ausschnittteils (8) freigibt, wobei das Band (14) auch wenigstens einen Schlitz (16) in der Nähe des Lochs (15) umfaßt, um so wenigstens eine Entlüftungsöffnung bereitzustellen.

2. Vorrichtung nach Anspruch 1, wobei das Band (14) Schlitz (16) an gegenüberliegenden Seiten des Lochs (15) umfaßt, um so zwei Entlüftungsöffnungen bereitzustellen.
3. Vorrichtung nach Anspruch 1, wobei der Ausschnittteil (8) 4 Millimeter auf 4,2 Millimeter ist.
4. Vorrichtung nach Anspruch 3, wobei das Verteilungsnetz (13) 6 Millimeter auf 5,8 Millimeter ist.
5. Vorrichtung nach Anspruch 4, wobei das Verteilungsnetz (13) mit Dioctylnatriumsulfosuccinat imprägniert ist.
6. Vorrichtung nach Anspruch 5, wobei das Loch (15) im Band (14) im wesentlichen den gesamten Ausschnittteil (8) freigibt.
7. Vorrichtung nach Anspruch 1 oder 6, des weiteren umfassend ein Strommeßgerät in elektrischer Verbindung mit der Arbeits- und Gegenelektrode.

### Revendications

1. Dispositif (1) pour détecter ou mesurer la concentration d'un analyte comprenant :

un premier isolateur électrique (2),  
une paire d'électrodes (4, 5) consistant en une électrode de travail et une contre-électrode, substantiellement de la même taille, les électrodes (4, 5) étant constituées des mêmes matériaux électriquement conducteurs et étant supportées par le premier isolateur électrique (2),  
un second isolateur électrique (3), recouvrant le premier isolateur électrique (2) et les électrodes (4, 5) et comportant une partie découpée (8),  
un réactif (11) pour détecter ou mesurer la concentration de l'analyte, le réactif (11) couvrant substantiellement les surfaces d'électrodes exposées dans la partie découpée (8) et étant en quantité suffisante pour analyser l'analyte dans un échantillon de sang complet d'au moins environ 9 microlitres et  
un tissu maillé diffuseur (13) recouvrant la partie découpée (8) et attaché au second isolateur électrique (3),

### caractérisé en ce que

la partie découpée (8) expose une surface plus petite de la contre-électrode que de l'électrode de travail, le tissu maillé diffuseur (13) est imprégné d'un surfactant,  
la partie découpée (8) et le tissu maillé diffuseur (13) sont d'une grandeur suffisante pour recevoir un échantillon de sang complet d'au moins environ 9 microlitres pour analyser l'analyte et  
le tissu maillé diffuseur (13) est attaché au second isolateur électrique (3) par un ruban (14) comportant un adhésif sur un côté et un trou (15) qui expose au moins une partie du tissu maillé diffuseur (13) et de la partie découpée (8) et dans lequel le ruban (14) comporte au moins une fente (16) près du trou (15) créant de cette façon au moins une prise d'air.

2. Dispositif selon la revendication 1, dans lequel le ruban (14) comporte des fentes (16) sur les côtés opposés au trou (15), créant de cette façon deux prises d'air.
3. Dispositif selon la revendication 1, dans lequel la partie découpée (8) mesure 4 millimètres sur 4,2 millimètres.
4. Dispositif selon la revendication 3, dans lequel le tissu maillé diffuseur (13) mesure 6 millimètres sur 5,8 millimètres.

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5. Dispositif selon la revendication 4, dans lequel le tissu maillé diffuseur (13) est imprégné de sulfosuccinate de dioctyle sodique.
6. Dispositif selon la revendication 5, dans lequel le trou (15) dans le ruban (14) expose substantiellement l'ensemble de la partie découpée (8).
7. Dispositif selon la revendication 1 ou 6 comprenant en outre un appareil de mesure du courant en liaison électrique avec l'électrode de travail et la contre-électrode.

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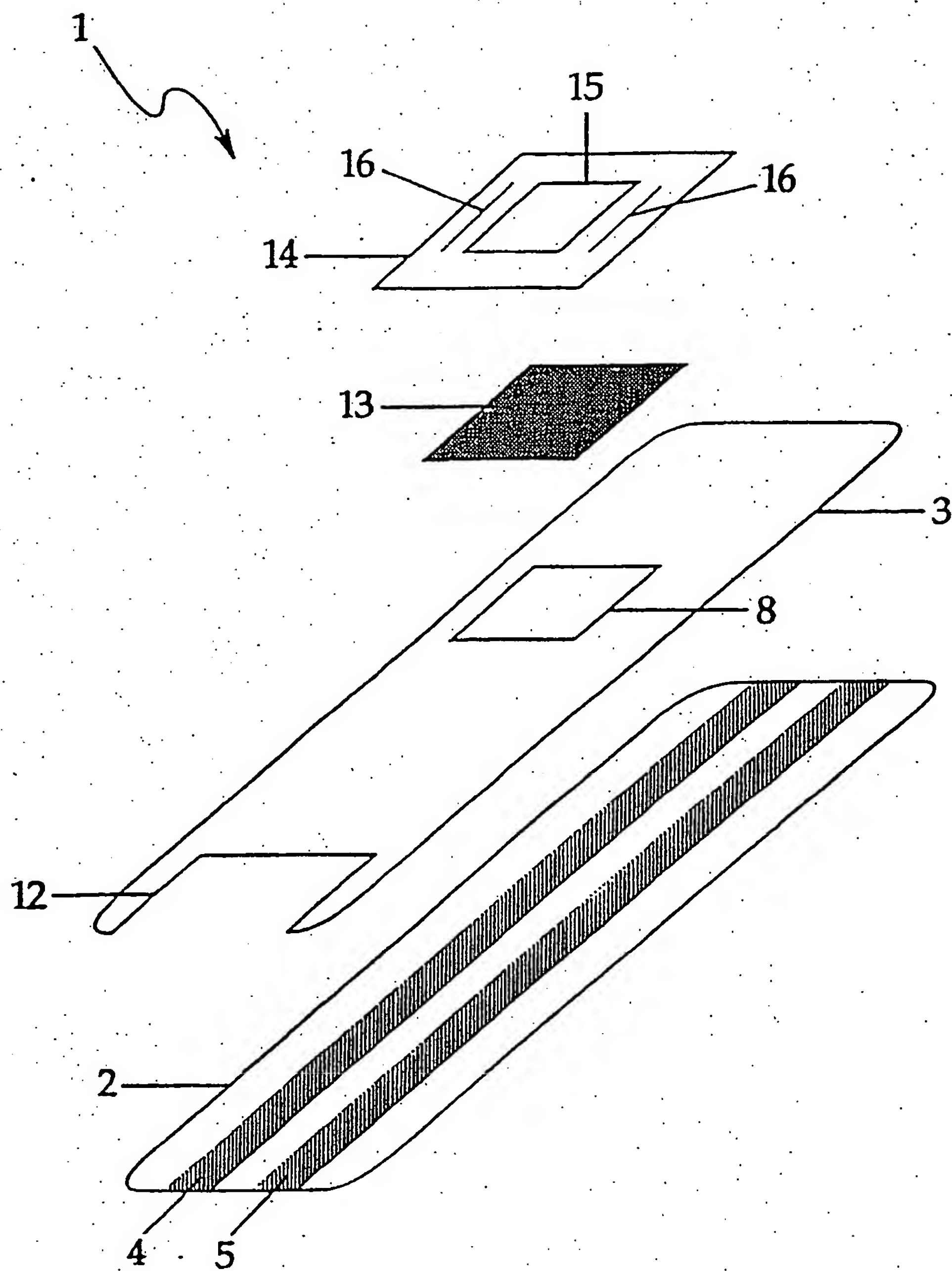


Fig. 1



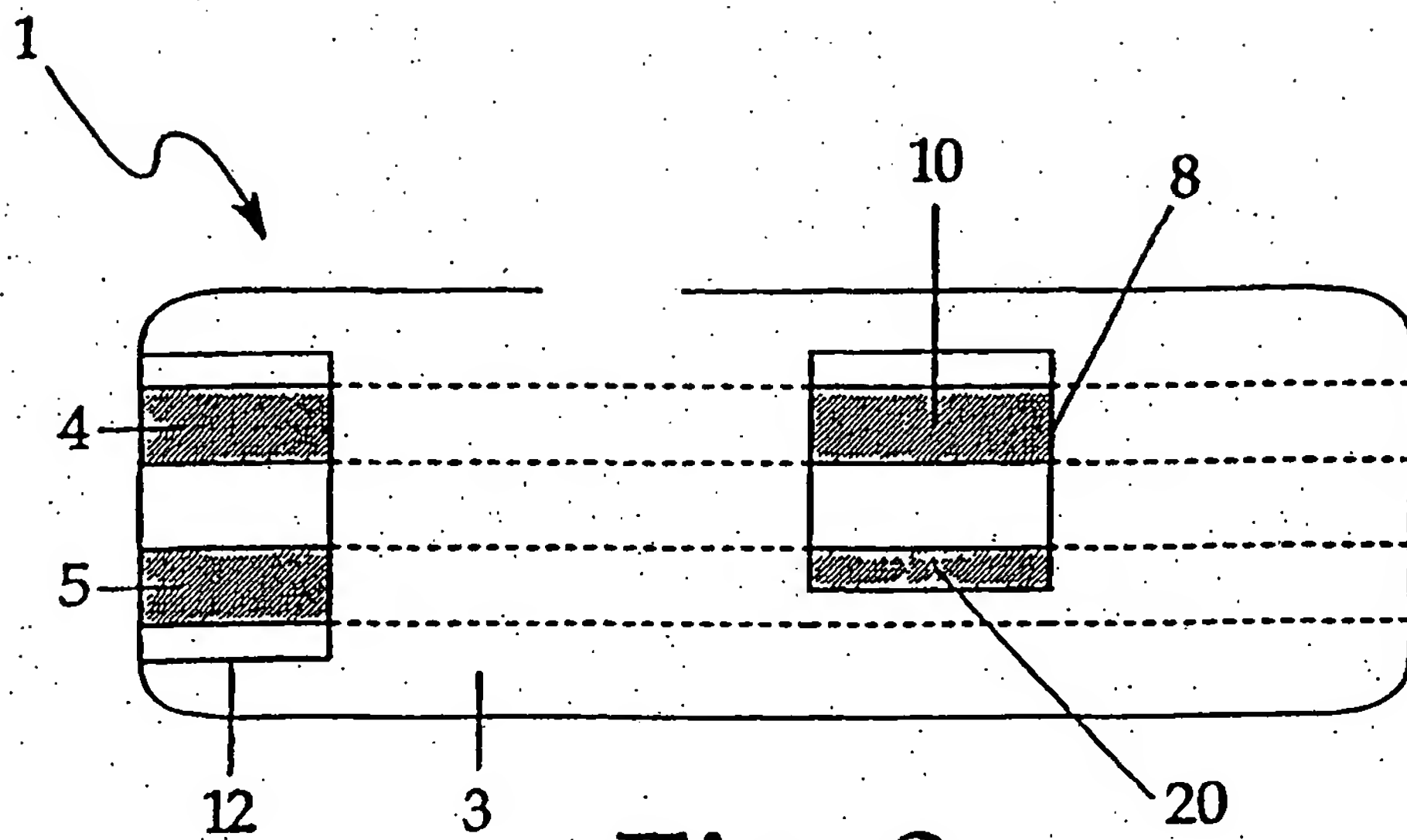


Fig. 2

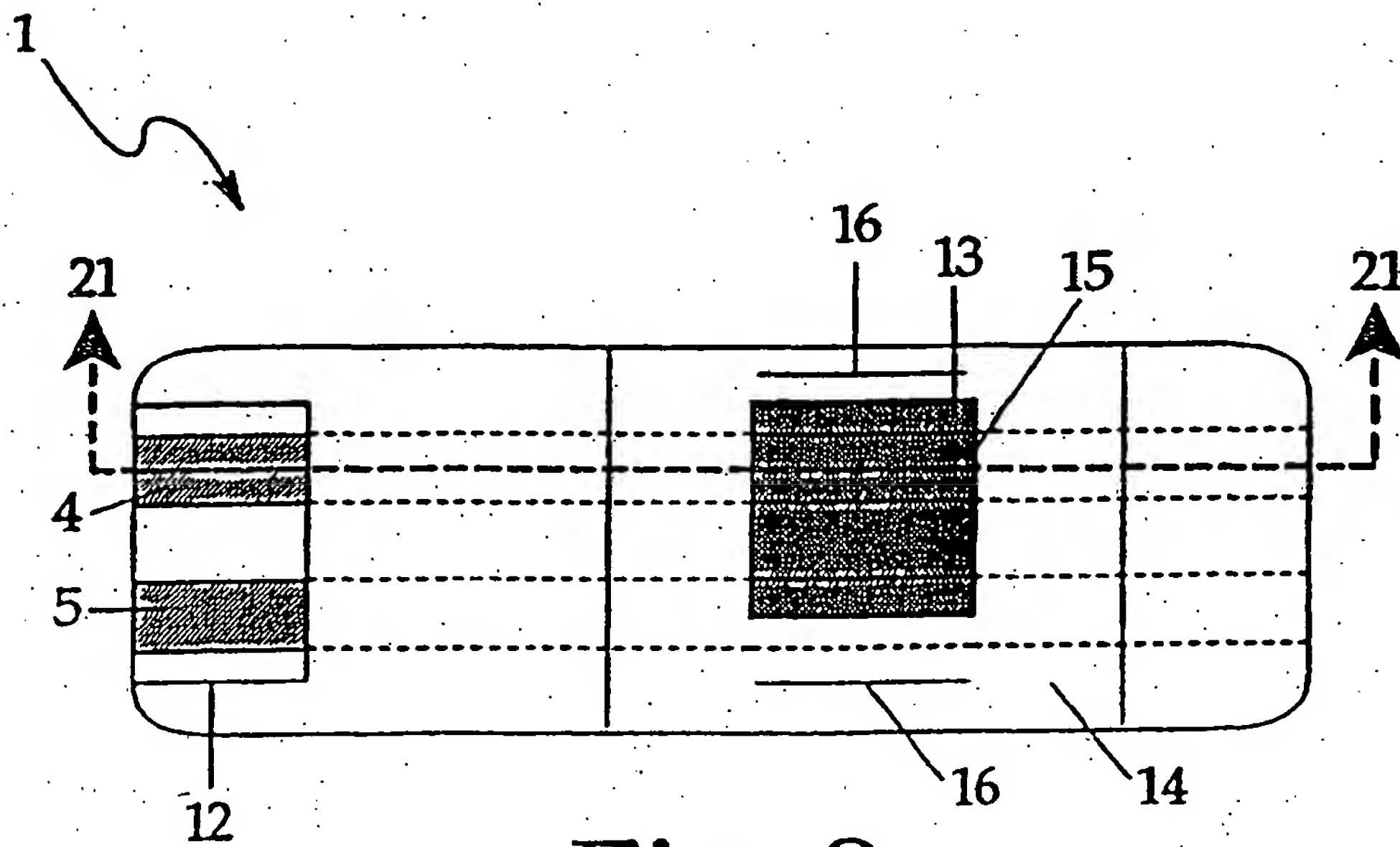


Fig. 3

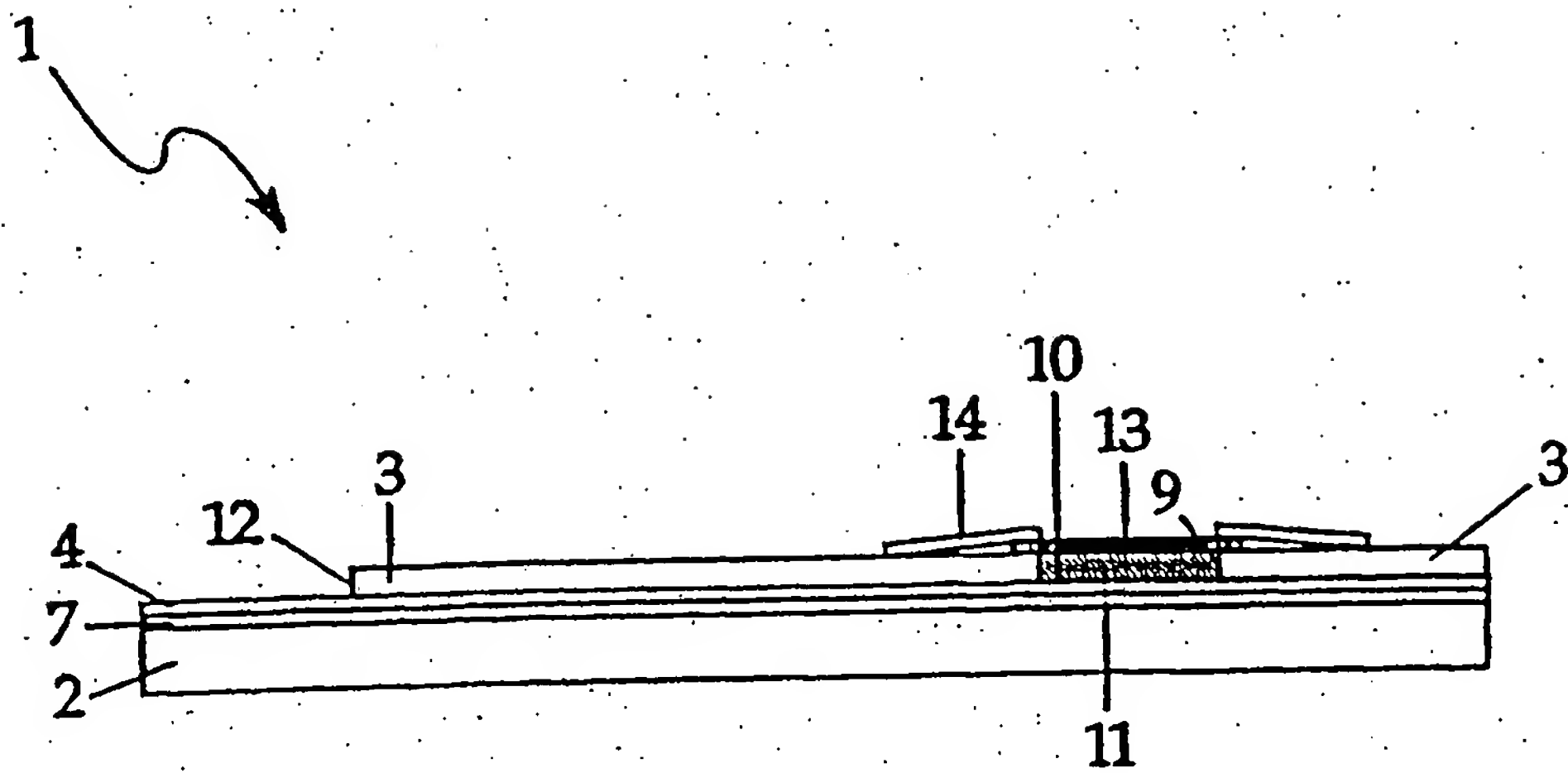


Fig. 4

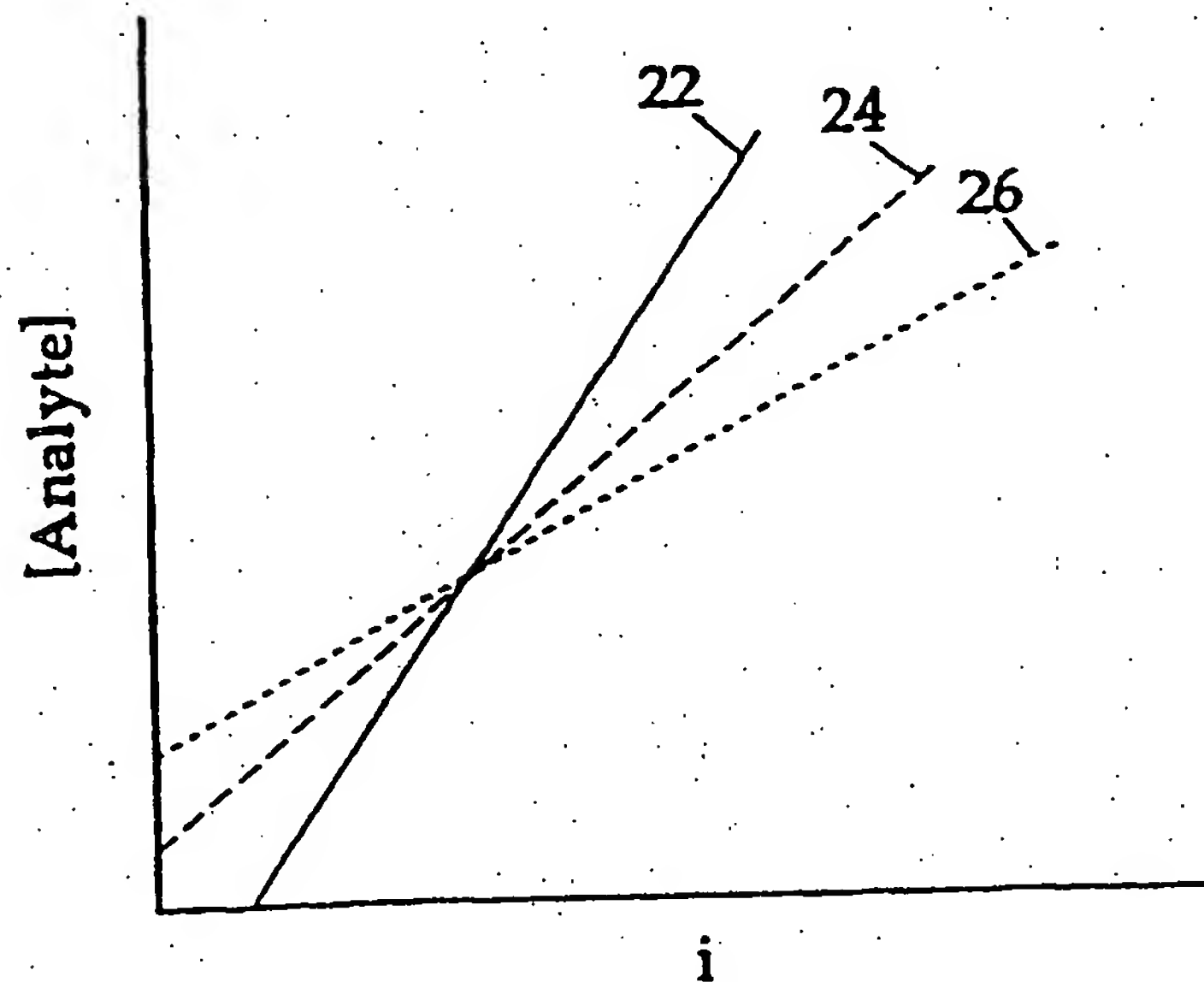


Fig. 5